## Amendments to the Claims:

The listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims**:

- 1. (Previously presented) Method for the identification of biomolecules in variant libraries of biomolecules comprising the steps:
  - a) Production of a variant library, consisting of a number of variants  $(B_0)$  of gene sequences coding for the biomolecule, and
  - b) Division of the variant library into a number of compartments  $(W_0)$ , which is at least by a factor of ten smaller than the number of variants in the variant library  $(B_0)$ , where each compartment contains a partial library which contains  $K_0=B_0/W_0$  variants,
  - c) Production of biomolecules in the compartments and testing of the biomolecules obtained in the single compartments for a specified phenotype, whereas from the observed phenotype no direct conclusions on the genotype can be made,
  - d) Selection of at least one compartment, which contains biomolecules fulfilling the wanted properties,
  - e) Division of the partial library contained in the selected compartment into further compartments, and
  - f) n-fold repetition of the steps c) to e) until in every compartment maximally only one variant  $(K_n \le 1)$  of the gene sequence coding for the biomolecule is contained.
  - 2. (Original) The method of claim 1, wherein the wanted property is a biocatalytic activity.

- 3. (Previously presented) The method of claim 1, wherein in step c) also an amplification of the partial library takes place in the compartments up to an number of individuals  $V_0(x)$  at the point in time x per compartment, whereas the number of individuals  $V_0(x)$  divided by the number of clones per compartment  $K_0$  gives the amplification factor  $F_0(x)$  per clone.
- 4. (Previously presented) The method of claim 1, wherein in step e) the division is carried out under dilution of the partial library by means of factor  $F_0(x)$ , so that in a given volume every clone contained in the compartment is statistically present up to a number  $X_0 < W_1$ , this volume is divided up in a number of new compartments  $W_1$ , whereas the new number of clones per compartment amounts to  $K_1 = X_0 * K_0 / W_1$ .
- 5. (Previously presented) The method of claim 1, wherein the variant library contains 103 to 10<sup>15</sup> variants of the gene sequence of the biomolecule.
- 6. (Previously presented) The method of claim 1, wherein in step b) the variant library is divided up in  $10^1$  to  $10^4$  compartments.
- 7. (Previously presented) The method of claim 1, wherein in step b) the variant library is transferred into an organism before division.
- 8. (Currently amended) The method of claim 7, wherein in step c) the culture of the organism after division is amplified to a number of organisms of 10<sup>8</sup> to 10<sup>9</sup> per compartment.
- 9. (Previously presented) The method of claim 7, wherein the organisms also conduct the production of the biomolecules.

- 10. (Previously presented) The method of claim 7, wherein the partial libraries in the compartments are re-isolated from the organisms, and the production of the biomolecules is conducted by cell-free systems.
- 11. (Currently amended) The method of claim [[1]] 3, wherein the amplification of the partial libraries and the production of the biomolecules is conducted by cell-free systems.
- 12. (Previously presented) The method of claim 1, wherein the variant library consists of DNA-plasmids, which contain the gene sequence coding for the biomolecule.
- 13. (Previously presented) The method of claim 1, wherein the variant library consists of linear nucleic acid molecules, which contain the gene sequence coding for the biomolecule.
- 14. (Previously presented) The method of claim 1, wherein the biomolecules are enzymes or ribozymes or other biomolecules, which exhibit a biocatalytic activity.
- 15. (Currently amended) The method of claim [[1]] 2, wherein the test for a biocatalytic activity is conducted with a physical detection method selected from the group consisting of UVIVIS-spectroscopy, fluorescence spectroscopy and fluorescence-correlation-spectroscopy.